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The effect of neuromuscular blockade on oxygen consumption in sedated and mechanically ventilated pediatric patients after cardiac surgery

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Abstract *Objective:* To measure the effect of intense neuromuscular blockade (NMB) on oxygen consumption (VO_2) in deeply sedated and mechanically ventilated children on the first day after complex congenital cardiac surgery. *Design:* Prospective clinical interventional study. *Setting:* Pediatric intensive care unit of an university medical centre. *Measurements and results:* Nine mechanically ventilated and sedated children (weight 2.8–8.7 kg) were included. All children were treated with vasoactive drugs. The level of sedation was quantified using the comfort score, Ramsay score and bispectral index (BIS). The intensity of NMB was quantified using acceleromyography and VO_2 was measured using indirect calorimetry. Analgo-sedation using various intravenous agents was targeted at a deep level (comfort score < 18, BIS < 60 and Ramsay score > 4). NMB was achieved by intravenous administration of rocuronium. All measurements were conducted before, during and after recovery from a period of intense NMB. Baseline values were VO_2 6.1 ml/(kg min) (SD 1.3), comfort score 13 (SD 0.7), BIS 42.5 (SD 14.2), mean blood pressure 54.0 mmHg (SD 10.5), mean heart rate 129.9 bpm (SD 28.9) and mean core temperature 36.7°C (SD 0.5). There were no significant differences in VO_2 or other parameters between baseline, during

NMB and the recovery phase.

Conclusion: Neuromuscular blocking agents do not reduce oxygen consumption in deeply sedated and mechanically ventilated children after congenital cardiac surgery.

Keywords Congenital cardiac surgery · Oxygen consumption · Pediatric · Intensive care

Abbreviations

VO_2	Oxygen consumption (ml/kg/min)
NMB	Neuromuscular blockade
PEEP	Positive end expiratory pressure (cmH ₂ O)
TOF	Train-of-four (ratio)
T1	First response in the train-of-four (%)
TCNT	TOF count
FiO ₂	Inspiratory oxygen fraction
PA	Pulmonary atresia
PDA	Patent ductus arteriosus
TPGV	Transposition great vessels
Fallot	Tetralogy of Fallot
AVSD	Atrioventricular septal defect
VSD	Ventricular septal defect
MAPCA	Major aorto-pulmonary collateral arteries
LOS	Length of stay
BIS	Bispectral index
SaO ₂	Arterial oxygen saturation (%)
MAP	Mean arterial pressure (mmHg)

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Introduction

Following complex congenital cardiac surgery, a certain number of patients suffer from a low cardiac output syndrome leading to a distorted balance between oxygen delivery (DO_2) and oxygen consumption (VO_2) [1]. Besides sedation and mechanical ventilation, the use of neuromuscular blocking agents has been recommended in order to decrease VO_2 [2].

To date, studies investigating the effect of neuromuscular blockade (NMB) on oxygen consumption show conflicting results, which may be explained by differences in study design [3–6].

We measured the effect of intense neuromuscular blockade on oxygen consumption in deeply sedated and mechanically ventilated children on the first day after complex congenital cardiac surgery.

Methods

This study was conducted in accordance with the Dutch national legislation concerning biomedical studies in children and was approved by the national and local ethics committees concerning medical research in humans. Oral and written informed consent was obtained from both parents.

We included children below 2 years of age on the first postoperative day following complex congenital cardiac surgery necessitating sedation and mechanical ventilation. Additional inclusion criteria were a body temperature between 36.0 and 38.0°C, FiO_2 less than 0.6 and air leak less than 10%. Exclusion criteria were severe renal insufficiency, allergy to rocuronium, and the use of NMB on the same day prior to the study. Hemodynamic and respiratory monitoring was routinely performed using a

HP Merlin monitoring system (HP, Irvine, USA). A rapidly changing blood pressure and/or heart rate were additional exclusion criteria.

All patients received analgo-sedation at the discretion of the treating physician (Table 1). Also, all children received inotropic agents consisting of a combination of dobutamine (mean dose 4.9 $\mu\text{g/kg/min}$) and/or nor-epinephrine (mean dose 0.4 $\mu\text{g/kg/min}$) combined with either enoximone or milrinone. Nutritional demands were met using either enteral feeding or intravenous dextrose as judged clinically necessary.

Oxygen consumption and carbon dioxide production were measured using the Deltatrac II metabolic monitor (GE healthcare, Chalfont St Giles, UK) as described previously [7]. This device provides a minute to minute VO_2 value.

The bispectral index monitor (BIS, Aspect Medical Systems, USA) is based upon a one channel EEG recording at the frontal location. The device provides a value between 0 and 100. A level of 100 equals a fully awake state, while a level between 40 and 60 reflect deep sedation or anesthesia. In children the BIS value correlates well with the level of sedation [8]. For clinical evaluation of the level of sedation the comfort score (CS) was used which has been described elsewhere [9, 10]. A score below 17 points reflects deep sedation. Although not validated in children we also determined the Ramsay Score (RS) [11]. The sedation scores were always determined by one author (JL) at the beginning, half-way and at the end of a recording period.

The intensity of NMB was guided by acceleromyographic recording of the thumb adduction following train-of-four stimulation (TOF) of the ulnar nerve with a 1-min interval (TOF-watch, Organon, Bostel, the Netherlands). This device has been validated in small children [12, 13].

Table 1 Patient data ($n = 9$)

Patient	Age (month)	Weight (kg)	Diagnosis	LOS PICU (days)	Mortality probability PIM (%)	Mortality probability PRISM II (%)	FiO_2 (fraction)	Analgo-sedation
1	7	4.9	PA, VSD, MAPCA's	4	1.4	1.8	0.40	mf, a, m, p, l
2	2	3	PDA, VSD	11	0.8	33.8	0.40	mf, a, m, p
3	17	8.7	TPGV + situs inversus	13	16.2	5.7	0.35	mf, a, m, p, l
4	6.5	7	Fallot + MAPCA's	14	4.8	5.9	0.60	mf, a, m, p
5	8	5.3	Fallot	7	2.3	5.9	0.60	mf, a, m, p, k
6	0.5	2.8	AVSD	18	4.6	8.9	0.31	mf, a, m
7	0.1	3	Hypoplastic right ventricle	4	3.0	8.9	0.21	mf, a, m, p
8	0.25	3.3	Intracardiac tumor	4	13.7	6.1	0.45	mf, a, m
9	6	6.4	VSD	10	0.8	5.9	0.60	mf, a, m, p, k
Mean value	5.3	4.9		9.4	5.2	9.2	0.40	

PA pulmonary atresia, MAPCA major aorto-pulmonary collateral arteries, PDA patent ductus arteriosus, TPGV transposition great vessels, Fallot fallot tetralogy, AVSD atrioventricular septal defect, VSD ventricular septal defect, LOS length of stay at the PICU, PIM

pediatric index of mortality, PRISMII pediatric risk of mortality version II, mf morphine, a acetaminophen, m midazolam, p propofol, l levopromazine, k s-ketamine

A TOF count (TCNT) of zero was considered intense NMB. A TOF ratio > 0.8 and the first response in the train-of-four (T_1) $> 80\%$ indicated sufficient clinical recovery from NMB.

We also collected PIM and PRISMII scores reflecting severity of illness [14, 15]. Furthermore, we collected data concerning length of stay (LOS), diagnosis, weight, height, and age. Mean blood pressure, heart rate, and core temperature were continuously measured.

At the start of the experiment, care was taken to provide a deep level of sedation with a CS less than or equal to 17, a RS higher than 4 and a BIS value less than 60. When this level of sedation was reached (but before NMB was instituted) a baseline registration was performed. Hereafter intense NMB was achieved by administering rocuronium in a dose of 0.3 mg/kg. If the TCNT did not decrease to 0 additional doses of rocuronium 0.15 mg/kg were given until the required TCNT was reached. At that point the registration of the NMB phase was started. During this phase, additional rocuronium (0.15 mg/kg) was administered if the TCNT value increased > 0 . After recording of the NMB phase time was taken to let the muscle relaxant wear off. When the TOF value exceeded 0.8 and $T_1 > 80\%$ recording of the recovery phase was performed.

A registration period in which all variables were collected lasted 20 min. The mean value of these 20 min was taken for further calculation.

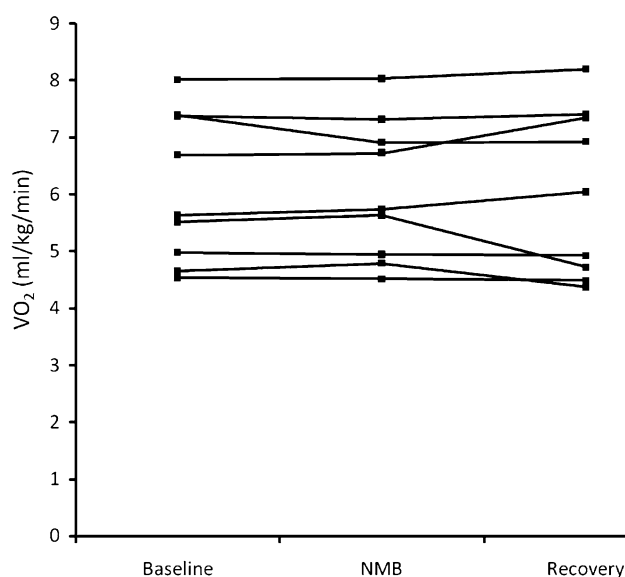
The coefficient of variation of the VO_2 measurement was calculated by dividing the standard deviation of measurements with 1-min interval by the mean value of the 20-min episode. Sample size calculation was performed using an expected variation of VO_2 measurement of 7%, a clinically significant change in VO_2 of at least 5%, a power of 80% and P value of 0.05. This resulted in a sample size of 13. Potential differences between various parameters during baseline, NMB and recovery were analyzed using ANOVA. Statistical calculations were performed using MedCalc version 9 (MedCalc Software, Mariakerke, Belgium).

Results

The coefficient of variation of the VO_2 measurements was 3.7% at baseline, 3.3% during NMB and 3.5% after recovery from NMB (mean 3.5%). Because the coefficient of variation was lower than anticipated (3.6 vs. 7%), we stopped the study after the inclusion of nine patients while maintaining the previously mentioned power.

Patient details are depicted in Table 1. The mean dose of rocuronium needed to achieve intense NMB was 0.7 mg/kg (range 0.3–1.2 mg/kg).

Figure 1 shows the individual changes in VO_2 for the nine patients during the experiment. Table 2 shows the



NMB = neuromuscular blockade

Fig. 1 Individual course of oxygen consumption during the baseline, NMB and recovery phase ($n = 9$)

mean values for several parameters before, during, and after neuromuscular blockade. Using ANOVA no changes in VO_2 or any of the other parameters were detected between baseline, during neuromuscular blockade, and the recovery phase except for intensity of NMB.

Discussion

This study shows that neuromuscular blocking agents do not reduce metabolic needs in deeply sedated and mechanically ventilated children after cardiac surgery.

In critically ill children one other study also observed a lack of difference between children with or without NMB [16]. However, most studies found a reduction in VO_2 after application of NMB [3–6, 17], although in some studies the effect was smaller or absent in patients without spontaneous movements before NMB was instituted [6]. One study in deeply sedated adults found similar results compared to our study [18]. We believe the level of sedation explains these differences. All children in our study were on controlled mechanical ventilation and had a documented deep level of sedation. Since NMB only affects skeletal muscle activity it seems logical to assume that, when muscle activity is already very low, a further reduction in VO_2 is not to be expected.

We made several precautions to optimize the validity of the recorded data. All measurements were recorded automatically; sedation scores were recorded by the same person and were supplemented by a measured BIS value.

Table 2 Mean values (SD) of various parameters before, during and after neuromuscular blockade ($n = 9$)

	Mean (SD)		
	Baseline	NMB	Recovery
VO ₂ (ml/kg/min)	6.1 (1.3)	6.1 (1.2)	6.1 (1.5)
BIS	42.5 (14.2)	42.0 (16.0)	46.1 (20.2)
Comfort score	13.0 (0.7)		12.3 (1.7)
Ramsay score	5.5 (0.5)		5.5 (0.5)
SaO ₂ (%)	94.4 (4.5)	93.9 (5.3)	94.3 (4.4)
Heart rate (bpm)	129.9 (28.9)	132.9 (30.7)	132.8 (28.5)
MAP (mmHg)	54.0 (10.5)	56.1 (13.5)	53.2 (8.7)
Core temperature (°C)	36.7 (0.5)	36.8 (0.5)	36.7 (0.6)
TI (%)	92.2 (10.1)	3.2 (2.7)*	95.1 (25.0)
TOF	1.14 (0.06)	0.00 (0.00)*	0.96 (0.07)

VO₂ oxygen consumption, BIS bispectral index, SaO₂ arterial oxygen saturation, MAP mean arterial pressure, TI relative force of first twitch, TOF train-of-four ratio

* Different from baseline and recovery; $P < 0.001$ (ANOVA)

Also we observed a small variation in the VO₂ measurement. Furthermore the intensity of the neuromuscular blockade was recorded using an established technique validated in (young) children. Using this technique we could actually demonstrate an intense level of NMB during the study and on the other hand a sufficient level of recovery at the end of the study.

The study is limited by several factors. With oxygen fractions in excess of 0.4 the accuracy of VO₂ measurement may decrease [7]. Although three children in our study were ventilated with a FiO₂ in excess of 0.4 (Table 1) the goal of this study was to detect changes in VO₂ within a relatively small time interval and because each patient acted as his or her own control we consider this to be of minor influence. The reliability of BIS and sedation scores might be questionable. It is therefore possible that several children reached a sedation level

which was equivalent to an anesthesia level. Caution is advised to extrapolate these results to other critically ill children, for example, with sepsis, trauma, or respiratory failure. Furthermore in our study only one child had a SaO₂ below 90% and our results may therefore not apply to children with a severe cyanosis. Lastly we only measured the effect of NMB on the first postoperative day. Results could differ when NMB is instituted in a later clinical phase.

While the level of sedation seems to be very important in the reduction of VO₂, the use of high dose sedatives seems attractive with the need for a reduction in VO₂. However these agents are associated with many side effects or complications [19] and might even reduce cardiac output. On the other hand, prolonged NMB is also associated with various possible side effects. The most feared complication is the critical illness neuromuscular disease [20].

Conclusion

Neuromuscular blocking agents do not reduce oxygen consumption and therefore do not reduce energy expenditure in well-sedated children following complex congenital cardiac surgery.

Conflict of interest statement None.

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